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INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification 6 :	A2	(11) International Publication Number: WO 99/65449
(52) International Application Number: PCT/US99/13632		(43) International Publication Date: 23 December 1999 (23.12.99)
(22) International Filing Date: 18 June 1999 (18.06.99)		(81) Designated States: AE, AL, AU, BA, BB, BG, BR, CA, CN, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, FR, GB, GR, GU, HK, IL, IN, IS, JP, KR, LC, LK, LU, LV, MG, MK, MN, MX, NO, NZ, PL, RO, SG, SI, SK, SL, TR, TT, UA, US, UZ, VN, YU, ZA, ZW)
(30) Priority Data: 60089,837	19 June 1998 (19.06.98)	US
(71) Applicant (for all designated States except US): SMITHKLINE BEECHAM CORPORATION [GB/GB]; One Franklin Plaza, Philadelphia, PA 19103 (GB).		
(72) Inventors; and		
(73) Inventors/Applicants (for US only): CALLAHAN, James, F. (US/US); 4214 Leones Street, Philadelphia, PA 19111 (US); CHAROT-PLETCHER, Marie, C. (US/US); 902 Gay Street, Phoenixville, PA 19460 (US).		
(74) Agents: STERGIO, Yuriy, P. et al.; SmithKline Beecham Corporation, Corporate Intellectual Property, UW2220, 709 Swedeland Road, P.O. Box 1339, King of Prussia, PA 19406-0939 (US).		

Without international search report and to be republished upon receipt of that report.

Published

(54) Title: INHIBITORS OF TRANSCRIPTION FACTOR NF- $\kappa$ B

(57) Abstract

The present invention provides pharmaceutical compositions of salicylanilide inhibitors of transcription factor NF- $\kappa$ B, and methods for treating diseases in which activation of NF- $\kappa$ B is implicated. More specifically, the present invention provides methods of treatment of a variety of diseases associated with NF- $\kappa$ B activation including inflammatory disorders; particularly rheumatoid arthritis, inflammatory bowel disease, and asthma; dermatitis, including psoriasis and atopic dermatitis; autoimmune diseases; tissue and organ rejection; Alzheimer's disease; stroke; atherosclerosis; restenosis; cancer, including Hodgkin's disease; certain viral infections, including AIDS; osteoarthritis; osteoporosis; and Axial Sclerostasis by administering to a patient in need thereof a compound of the present invention.

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# **INHIBITORS OF TRANSCRIPTION FACTOR NF- $\kappa$ B**

## **FIELD OF THE INVENTION**

This invention relates in general to salicylanilide inhibitors of transcription factor NF- $\kappa$ B. Such compounds are particularly useful for treating diseases in which activation of NF- $\kappa$ B is implicated. More specifically, these compounds inhibit I $\kappa$ B phosphorylation and subsequent degradation. Such compounds are useful in the treatment of a variety of diseases associated with NF- $\kappa$ B activation including inflammatory disorders, particularly rheumatoid arthritis, inflammatory bowel disease, and asthma; dermatosis, including psoriasis and atopic dermatitis; autoimmune diseases; tissue and organ rejection; Alzheimer's disease; stroke; atherosclerosis; restenosis; cancer, including Hodgkins disease; and certain viral infections, including AIDS; osteoarthritis; osteoporosis; and Anaxia Telangiectasia.

## **BACKGROUND OF THE INVENTION**

Recent advances in scientific understanding of the mediators involved in acute and chronic inflammatory diseases and cancer have led to new strategies in the search for effective therapeutics. Traditional approaches include direct target intervention such as the use of specific antibodies, receptor antagonists, or enzymic inhibitors. Recent breakthroughs in the elucidation of regulatory mechanisms involved in the transcription and translation of a variety of mediators have led to increased interest in therapeutic approaches directed at the level of gene transcription.

NF- $\kappa$ B belongs to a family of closely related dimeric transcription factor complexes composed of various combinations of the Rel/NF- $\kappa$ B family of polypeptides. The family consists of five individual gene products in mammals, RelA (p65), NF- $\kappa$ B1 (p50/p105), NF- $\kappa$ B2 (p49/p100), c-Rel, and RelB, all of which can form hetero- or homodimers. These proteins share a highly homologous 300 amino acid "Rel homology domain" which contains the DNA binding and dimerization domains. At the extreme C-terminus of the Rel homology domain is a nuclear translocation sequence important in the transport of NF- $\kappa$ B from the cytoplasm to the nucleus. In addition, p65 and cRel possess potent transactivation domains at their C-terminal ends.

The activity of NF- $\kappa$ B is regulated by its interaction with a member of the inhibitor I $\kappa$ B family of proteins. This interaction effectively blocks the nuclear localization sequence

on the NF- $\kappa$ B proteins, thus preventing migration of the dimer to the nucleus. A wide variety of stimuli activate NF- $\kappa$ B through what are likely to be multiple signal transduction pathways. Included are bacterial products (LPS), some viruses (HIV-1, HTLV-1), inflammatory cytokines (TNF $\alpha$ , IL-1), and environmental stress. Apparently common to all stimuli however, is the phosphorylation and subsequent degradation of I $\kappa$ B. I $\kappa$ B is phosphorylated on two N-terminal serines by the recently identified I $\kappa$ B kinases (IKK- $\alpha$  and IKK- $\beta$ ). Site-directed mutagenesis studies indicate that these phosphorylations are critical for the subsequent activation of NF- $\kappa$ B in that once phosphorylated the protein is flagged for degradation via the ubiquitin-proteasome pathway. Free from I $\kappa$ B, the active NF- $\kappa$ B complexes are able to translocate to the nucleus where they bind in a selective manner to preferred gene-specific enhancer sequences. Included in the genes regulated by NF- $\kappa$ B are a number of cytokines, cell adhesion molecules, and acute phase proteins.

It is well-known that NF- $\kappa$ B plays a key role in the regulated expression of a large number of pro-inflammatory mediators including cytokines such as IL-6 and IL-8, cell adhesion molecules, such as ICAM and VCAM, and inducible nitric oxide synthase (iNOS). Such mediators are known to play a role in the recruitment of leukocytes at sites of inflammation and in the case of iNOS, may lead to organ destruction in some inflammatory and autoimmune diseases.

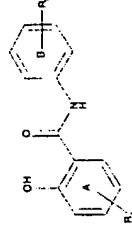
The importance of NF- $\kappa$ B in inflammatory disorders is further strengthened by studies of airway inflammation including asthma, in which NF- $\kappa$ B has been shown to be activated. This activation may underlie the increased cytokine production and leukocyte infiltration characteristic of these disorders. In addition, inhaled steroids are known to reduce airway hyperresponsiveness and suppress the inflammatory response in asthmatic airways. In light of the recent findings with regard to glucocorticoid inhibition of NF- $\kappa$ B, one may speculate that these effects are mediated through an inhibition of NF- $\kappa$ B.

Further evidence for a role of NF- $\kappa$ B in inflammatory disorders comes from studies of rheumatoid synovium. Although NF- $\kappa$ B is normally present as an inactive cytoplasmic complex, recent immunohistochemical studies have indicated that NF- $\kappa$ B is present in the nuclei, and hence active, in the cells comprising rheumatoid synovium. Furthermore, NF- $\kappa$ B has been shown to be activated in human synovial cells in response to stimulation with TNF- $\alpha$ . Such a distribution may be the underlying mechanism for the increased cytokine

psoriasis and atopic dermatitis; autoimmune diseases; tissue and organ rejection; Alzheimer's disease; stroke; atherosclerosis; testostosis; cancer, including Hodgkins disease; and certain viral infections, including AIDS; osteoarthritis; osteoporosis; and Ataxia Telangiectasia.

# DETAILED DESCRIPTION OF THE INVENTION

The present invention provides a method of treatment of diseases associated with NF- $\kappa$ B activation, comprising administering to an animal, particularly a mammal, most particularly a human in need thereof a compound of Formula I:

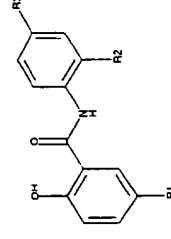


wherein:

R<sub>A</sub> substitutes ring A 0-3 times and is independently selected from the group consisting of: NO<sub>2</sub>, halogen, C<sub>1</sub>-alkyl, trifluoromethyl, O-C<sub>1</sub>-alkyl and S-C<sub>1</sub>-alkyl; and R<sub>B</sub> substitutes ring B 0-3 times and is independently selected from the group consisting of: halogen, C(O)C<sub>1</sub>-alkyl, C<sub>1</sub>-alkyl, O-C<sub>1</sub>-alkyl, S-C<sub>1</sub>-alkyl, CH<sub>2</sub>-aryl, and aryl;

and pharmaceutically acceptable salts, hydrates and solvates thereof.

The present invention further provides a preferred method of treatment of diseases associated with NF- $\kappa$ B activation, comprising administering to an animal, particularly a mammal, most particularly a human in need thereof a compound of Formula II:



II

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and eicosanoid production characteristic of this tissue. See Roshak, A. K., et al., *J. Biol. Chem.*, 271, 31496-31501 (1996).

The NF- $\kappa$ B/Rel and I $\kappa$ B proteins are also likely to play a key role in neoplastic transformation. Family members are associated with cell transformation *in vitro* and *in vivo* as a result of overexpression, gene amplification, gene rearrangements or translocations. In addition, rearrangement and/or amplification of the genes encoding these proteins are seen in 20-25% of certain human lymphoid tumors. In addition, a role for NF- $\kappa$ B in the regulation of apoptosis has been reported strengthening the role of this transcription factor in the control of cell proliferation.

Several NF- $\kappa$ B inhibitors are described in C. Wühl, et al., *J. Clin. Invest.* 101(5), 1163-1174 (1998), R. W. Sullivan, et al., *J. Med. Chem.* 41, 413-419 (1998), J. W. Pierce, et al., *J. Biol. Chem.* 272, 21096-21103 (1997)

The marine natural product hymenialdisine is known to inhibit NF- $\kappa$ B. Roshak, A., et al., *JPET*, 283, 955-961 (1997). Breton, J. J and Chabot-Fletcher, M. C., *JPET*, 282, 459-466 (1997).

Salicylanilides are known compounds. A general solution preparation of salicylanilides is described by M. T. Clark, R. A. Cuburn, R. T. Evans, R. J. Genco, *J. Med. Chem.*, 1986, 29, 25-29.

We have now discovered a novel method of inhibiting the activation transcription factor NF- $\kappa$ B using salicylanilides.

## SUMMARY OF THE INVENTION

An object of the present invention is to provide a method for treating diseases which may be therapeutically modified by altering the activity of transcription factor NF- $\kappa$ B.

Accordingly, in the first aspect, this invention provides a pharmaceutical composition comprising a compound according to Formula I.

In still another aspect, this invention provides a method of treating diseases in which the disease pathology may be therapeutically modified by inhibiting NF- $\kappa$ B.

In a particular aspect, this invention provides methods for treating a variety of diseases associated with NF- $\kappa$ B activation including inflammatory disorders; particularly rheumatoid arthritis, inflammatory bowel disease, and asthma; dermatosis, including

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wherein  $R_A$  of Formula I occurs once and is  $R_1$ ; and  $R_B$  of Formula I occurs twice and is independently  $R_2$  and  $R_3$ . More specifically:

$R_1$  is selected from the group consisting of: H,  $NO_2$ ,  $CF_3$ , F, Cl, Br, and I;

$R_2$  is selected from the group consisting of: H and F; and

$R_3$  is selected from the group consisting of: F, Cl, Br, I, phenyl and  $C(O)C_1$ -alkyl, preferably  $C_1$ -alkyl is  $CH_3$ ;

and pharmaceutically acceptable salts, hydrates and solvates thereof.

Compounds of Formula II selected from the following group are most preferred for use in the methods of the present invention:

$N-(4\text{-phenyl-phenyl})\text{-}2\text{-hydroxy-}5\text{-trifluoromethylcarboxamide}$ ;

$N-(2,4\text{-difluorophenyl})\text{-}2\text{-hydroxy-}5\text{-nitrocarboxamide}$ ;

$N-(2,4\text{-difluorophenyl})\text{-}2\text{-hydroxy-}5\text{-iodocarboxamide}$ ; and

$N-(4\text{-acetylphenyl})\text{-}2\text{-hydroxy-}5\text{-iodocarboxamide}$ .

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#### Definitions

The present invention includes all hydrates, solvates, complexes and prodrugs of the compounds of this invention. Prodrugs are any covalently bonded compounds which release the active parent drug according to Formulas I and II *in vivo*. If a chiral center or another form of an isomeric center is present in a compound of the present invention, all

forms of such isomer or isomers, including enantiomers and diastereomers, are intended to be covered herein. Inventive compounds containing a chiral center may be used as a racemic mixture, an enantiomerically enriched mixture, or the racemic mixture may be separated using well-known techniques and an individual enantiomer may be used alone. In cases in which compounds have unsaturated carbon-carbon double bonds, both the *cis*

(*Z*) and *trans* (*E*) isomers are within the scope of this invention. In cases wherein

compounds may exist in tautomeric forms, such as keto-enol tautomers, each tautomeric form is contemplated as being included within this invention whether existing in equilibrium or predominantly in one form.

The meaning of any substituent at any one occurrence in Formula I or any subformula thereof is independent of its meaning, or any other substituent's meaning, at any other occurrence, unless specified otherwise.

" $C_1$ -alkyl" as applied herein is meant to include substituted and unsubstituted methyl, ethyl, n-propyl, isopropyl, n-butyl, isobutyl and t-butyl, pentyl, n-pentyl, isopentyl,

neopentyl and hexyl and the simple aliphatic isomers thereof. Any  $C_1$ -alkyl group may be optionally substituted independently by one or two halogens, SR', OR',  $N(R')_2$ ,  $C(O)N(R')_2$ , carbamyl or  $C_1$ -4alkyl, where R' is  $C_1$ -alkyl.

"Halogen" as applied herein is meant to include F, Cl, Br, and I.

"Ar" or "aryl" as applied herein is meant to include phenyl or naphthyl, optionally substituted by one or more of  $Ph-C_0$ -alkyl;  $Het-C_0$ -alkyl;  $C_1$ -alkoxy;  $Ph-C_0$ -alkoxy;  $Het-C_0$ -alkoxy;  $OH$ ;  $(CH_2)_1-6NR^4R^5$ ;  $O(CH_2)_1-6NR^4R^5$ ;  $C_1$ -alkyl; OR';  $N(R')_2$ ; SR';  $CF_3$ ;  $NO_2$ ; CN;  $CO_2R'$ ;  $CON(R')$ ; F, Cl, Br or I; where  $R^4$  and  $R^5$  are H,  $C_1$ -alkyl,  $Ph-C_0$ -alkyl, or naphthyl- $C_0$ -alkyl; and R' is phenyl, naphthyl, or  $C_1$ -alkyl.

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#### Methods of Preparation

The compounds of the present invention may be conveniently prepared by the methods set forth in Schemes 1 & 2 below.

A general solution preparation of salicylanilides is described by M. T. Clark, R. A. Coburn, R. T. Evans, R. J. Genco; *J. Med. Chem.* 1986; 29: 25-29. These compounds may also be conveniently prepared by solid phase techniques, including in the form of a library.

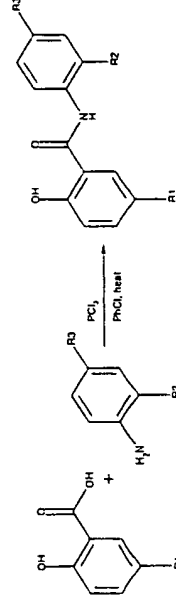
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#### General Preparation:

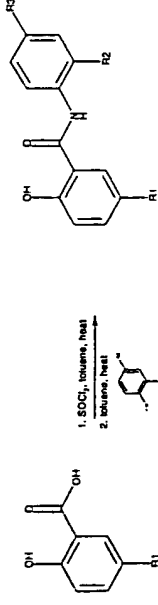
The general solution preparation is shown in Scheme 1 and 2. A salicylanilide is prepared by the condensation of a substituted salicylic acid with a substituted aniline in the presence of phosphorous trichloride in dry chlorobenzene (Scheme 1). A salicylanilide can also be prepared by conversion of a substituted salicylic acid to its corresponding acid chloride with thionyl chloride in toluene with catalytic DMF. The resulting salicylic acid chloride and a substituted aniline is then heated in toluene to form a salicylanilide (Scheme 2).

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Scheme 1



Scheme 2



Referring to the methods of preparing the compounds of Formula I set forth in

- 5 Schemes 1 & 2 above, the skilled artisan will appreciate that the present invention includes all novel intermediates required to make the compounds of Formula 1.

The starting materials used herein are commercially available or are prepared by routine methods well known to those of ordinary skill in the art and can be found in

standard reference books, such as the COMPENDIUM OF ORGANIC SYNTHETIC

- 10 METHODS, Vol. I-VI (published by Wiley-Interscience).

Acid addition salts of the compounds of Formula I are prepared in a standard

manner in a suitable solvent from the parent compound and an excess of an acid, such as

hydrochloric, hydrobromic, hydrofluoric, sulfuric, phosphoric, acetic, trifluoroacetic,

maleic, succinic or methanesulfonic. Certain of the compounds form inner salts or

zwitterions which may be acceptable. Cationic salts are prepared by treating the parent

compound with an excess of an alkaline reagent, such as a hydroxide, carbonate or

alkoxide, containing the appropriate cation; or with an appropriate organic amine. Cations

such as Li<sup>+</sup>, Na<sup>+</sup>, K<sup>+</sup>, Ca<sup>++</sup>, Mg<sup>++</sup> and NH<sub>4</sub><sup>+</sup> are specific examples of cations present in

pharmaceutically acceptable salts. Halides, sulfate, phosphate, alkanoates (such as acetate

and trifluoroacetate), benzoates, and sulfonates (such as mesylate) are examples of anions

present in pharmaceutically acceptable salts.

This invention provides a pharmaceutical composition which comprises a compound according to Formula I and a pharmaceutically acceptable carrier, diluent or

excipient. Accordingly, the compounds of Formula I may be used in the manufacture of a

medicament. Pharmaceutical compositions of the compounds of Formula I prepared as

hereinbefore described may be formulated as solutions or lyophilized powders for

parenteral administration. Powders may be reconstituted by addition of a suitable diluent

or other pharmaceutically acceptable carrier prior to use. The liquid formulation may be a

buffered, isotonic, aqueous solution. Examples of suitable diluents are normal isotonic

saline solution, standard 5% dextrose in water or buffered sodium or ammonium acetate solution. Such formulation is especially suitable for parenteral administration, but may also be used for oral administration or contained in a metered dose inhaler or nebulizer for insufflation. It may be desirable to add excipients such as polyvinylpyrrolidone, gelatin, hydroxy cellulose, acacia, polyethylene glycol, mannitol, sodium chloride or sodium citrate.

Alternately, these compounds may be encapsulated, tableted or prepared in an

emulsion or syrup for oral administration. Pharmaceutically acceptable solid or liquid

carriers may be added to enhance or stabilize the composition, or to facilitate preparation of

- 10 the composition. Solid carriers include starch, lactose, calcium sulfate dihydrate, terra alba,

magnesium stearate or stearic acid, talc, pectin, acacia, agar or gelatin. Liquid carriers

include syrup, peanut oil, olive oil, saline and water. The carrier may also include a

sustained release material such as glyceryl monostearate or glyceryl distearate, alone or

with a wax. The amount of solid carrier varies but, preferably, will be between about 20

- 15 mg to about 1 g per dosage unit. The pharmaceutical preparations are made following the

conventional techniques of pharmacy involving milling, mixing, granulating, and

compressing, when necessary, for tablet forms; or milling, mixing and filling for hard

gelatin capsule forms. When a liquid carrier is used, the preparation will be in the form of

a syrup, elixir, emulsion or an aqueous or non-aqueous suspension. Such a liquid

- 20 formulation may be administered directly p.o. or filled into a soft gelatin capsule.

For rectal administration, the compounds of this invention may also be combined

with excipients such as cocoa butter, glycerin, gelatin or polyethylene glycols and molded

into a suppository.

The methods of the present invention include topical administration of the

compounds of Formulas I and II. By topical administration is meant non-systemic

- 25 administration, including the application of a compound of the invention externally to the

epidermis, to the buccal cavity and instillation of such a compound into the ear, eye and

nose, wherein the compound does not significantly enter the blood stream. By systemic

administration is meant oral, intravenous, intraperitoneal and intramuscular administration.

- 30 The amount of a compound of the invention (hereinafter referred to as the active ingredient)

required for therapeutic or prophylactic effect upon topical administration will, of course,

vary with the compound chosen, the nature and severity of the condition being treated and

the animal undergoing treatment, and is ultimately at the discretion of the physician

While it is possible for an active ingredient to be administered alone as the raw chemical, it is preferable to present it as a pharmaceutical formulation. The active ingredient may comprise, for topical administration, from 0.01 to 5.0 wt% of the formulation.

The topical formulations of the present invention, both for veterinary and for human medical use, comprise an active ingredient together with one or more acceptable carriers therefor, and optionally any other therapeutic ingredients. The carrier must be "acceptable" in the sense of being compatible with the other ingredients of the formulation and not deleterious to the recipient thereof.

Formulations suitable for topical administration include liquid or semi-liquid preparations suitable for penetration through the skin to the site of where treatment is required such as: liniments, lotions, creams, ointments or pastes, and drops suitable for administration to the eye, ear or nose.

Drops according to the present invention may comprise sterile aqueous or oily solutions or suspensions and may be prepared by dissolving the active ingredient in a suitable aqueous solution of a bactericidal and/or fungicidal agent and/or any other suitable preservative, and preferably including a surface active agent. The resulting solution may then be clarified by filtration, transferred to a suitable container which is then sealed and sterilized by autoclaving or maintaining at 90-100 C for half an hour. Alternatively, the solution may be sterilized by filtration and transferred to the container by an aseptic technique. Examples of bactericidal and fungicidal agents suitable for inclusion in the drops are phenylmercuric nitrate or acetate (0.002%), benzalkonium chloride (0.01%) and chlorhexidine acetate (0.01%). Suitable solvents for the preparation of an oily solution include glycerol, diluted alcohol and propylene glycol.

Lotions according to the present invention include those suitable for application to the skin or eye. An eye lotion may comprise a sterile aqueous solution optionally containing a bactericide and may be prepared by methods similar to those for the preparation of drops. Lotions or liniments for application to the skin may also include an agent to hasten drying and to cool the skin, such as an alcohol or acetone, and/or a moisturizer such as glycerol or an oil such as castor oil or arachis oil.

Creams, ointments or pastes according to the present invention are semi-solid formulations of the active ingredient for external application. They may be made by mixing the active ingredient in finely-divided or powdered form, alone or in solution or

suspension in an aqueous or non-aqueous fluid, with the aid of suitable machinery, with a greasy or non-greasy basis. The basis may comprise hydrocarbons such as hard, soft or liquid paraffin, glycerol, beeswax, a mucilage, an oil of natural origin such as almond, corn, arachis, castor or olive oil; wool fat or its derivatives, or a fatty acid such as stearic or oleic acid together with an alcohol such as propylene glycol or macrogols. The formulation may incorporate any suitable surface active agent such as an anionic, cationic or non-ionic surface active agent such as sorbitan esters or polyoxyethylene derivatives thereof. Suspending agents such as natural gums, cellulose derivatives or in organic materials such as siliceous silicas, and other ingredients such as lanolin, may also be included.

# Utility of the Present Invention

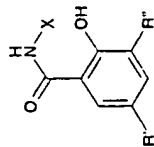
The compounds of Formulas I and II are useful as inhibitors of NF- $\kappa$ B. The present invention provides useful compositions and formulations of said compounds, including pharmaceutical compositions and formulations of said compounds.

The present invention also provides methods of treatment of diseases associated with NF- $\kappa$ B activation, which methods comprise administering to an animal, particularly a mammal, most particularly a human in need thereof a compound of Formulas I or II. The present invention particularly provides methods for treating inflammatory disorders; particularly rheumatoid arthritis, inflammatory bowel disease, and asthma; dermatosis, including psoriasis and atopic dermatitis; autoimmune diseases; tissue and organ rejection; Alzheimer's disease; stroke; atherosclerosis; restenosis; cancer, including Hodgkins disease; and certain viral infections, including AIDS; osteoarthritis; osteoporosis; and Ataxia Telangiectasia.

For acute therapy, parenteral administration of a compound of Formulas I or II is preferred. An intravenous infusion of the compound in 5% dextrose in water or normal saline, or a similar formulation with suitable excipients, is most effective, although an intramuscular bolus injection is also useful. Typically, the parenteral dose will be about 0.01 to about 50 mg/kg; preferably between 0.1 and 20 mg/kg, in a manner to maintain the concentration of drug in the plasma at a concentration effective to inhibit activation of NF- $\kappa$ B. The compounds are administered one to four times daily at a level to achieve a total daily dose of about 0.4 to about 80 mg/kg/day. The precise amount of an inventive compound which is therapeutically effective, and the route by which such compound is best

Table 1

Inhibition of NF- $\kappa$ B-driven Reporter Gene Activity



Compound Number	R'	R''	X	IC <sub>50</sub> $\mu$ M
1	H	H		8.14
2	H	H		~5
3	NO <sub>2</sub>	H		7.20
4	H	Ph		inactive
5	Ph	H		inactive
6	Br	H		0.82
7	CF <sub>3</sub>	H		36.5
8	CF <sub>3</sub>	H		3.55

NF- $\kappa$ B plays a key role in the regulated expression of a large number of pro-inflammatory mediators including cytokines such as IL-6 and IL-8 (Mukaida *et al.*, 1990; Liberman and Baltimore, 1990; Matsusaka *et al.*, 1993), cell adhesion molecules, such as ICAM and VCAM (Mani *et al.*, 1993; Kawai *et al.*, 1995; Ledebur and Parks, 1995), and inducible nitric oxide synthase (iNOS) (Xie *et al.*, 1994; Adcock *et al.*, 1994). (Full reference citations are at the end of this section). Such mediators are known to play a role in the recruitment of leukocytes at sites of inflammation and in the case of iNOS, may lead to organ destruction in some inflammatory and autoimmune diseases (McCarney-Francis *et al.*, 1993; Kleemann *et al.*, 1993). Importantly, the compounds described herein inhibit IL-8 synthesis and the production of nitric oxide, a product of iNOS activity (*see* Table 2).

administered, is readily determined by one of ordinary skill in the art by comparing the blood level of the agent to the concentration required to have a therapeutic effect.

The compounds of Formulas I and II may also be administered orally to the patient, in a manner such that the concentration of drug is sufficient to inhibit NF- $\kappa$ B or to achieve any other therapeutic indication as disclosed herein. Typically, a pharmaceutical composition containing the compound is administered at an oral dose of between about 0.1 to about 50 mg/kg in a manner consistent with the condition of the patient. Preferably the oral dose would be about 0.5 to about 20 mg/kg.

The compounds of Formulas I and II may also be administered topically to the patient, in a manner such that the concentration of drug is sufficient to inhibit NF- $\kappa$ B or to achieve any other therapeutic indication as disclosed herein. Typically, a pharmaceutical composition containing the compound is administered in a topical formulation of between about 0.01% to about 5% w/w.

No unacceptable toxicological effects are expected when compounds of the present invention are administered in accordance with the present invention.

The ability of the compounds described herein to inhibit the activation of NF- $\kappa$ B is clearly evidenced in their ability to inhibit NF- $\kappa$ B-driven reporter gene activity (*see* Table 1). The utility of the present NF- $\kappa$ B inhibitors in the therapy of diseases is premised on the importance of NF- $\kappa$ B activation in a variety of diseases.

TABLE 2

Anti-inflammatory Activity of Compound 3 in Table 1

<i>In vitro</i>	
TNF-stimulated IL-8 production in U937 cells	IC <sub>50</sub> = 3 uM
IL-1-induced PGE2 production in RSF	IC <sub>50</sub> = 3 uM
IL-1-stimulated Nitric Oxide production	IC <sub>50</sub> = 5 uM
<i>In vivo</i>	
Phorbol-ester-induced ear inflammation	ED <sub>50</sub> = 0.2 mg/ear
Ear swelling	ED <sub>50</sub> = 0.2 mg/ear
Inflammatory cell infiltration	

Evidence for an important role of NF-κB in inflammatory disorders is obtained in studies of asthmatic patients. Bronchial biopsies taken from mild atopic asthmatics show significant increases in the number of cells in the submucosa staining for activated NF-κB, total NF-κB, and NF-κB-regulated cytokines such as GM-CSF and TNFα compared to biopsies from normal non-atopic controls (Wilson *et al.*, 1998). Furthermore, the percentage of vessels expressing NF-κB immunoreactivity is increased as is IL-8 immunoreactivity in the epithelium of the biopsy specimens (Wilson *et al.*, 1998). As such, inhibition of IL-8 production through the inhibition of NF-κB, as has been demonstrated by these compounds would be predicted be beneficial in airway inflammation.

Recent studies suggest that NF-κB may also play a critical role in the pathogenesis of inflammatory bowel disease (IBD). Activated NF-κB is seen in colonic biopsy specimens from Chron's disease and ulcerative colitis patients (Ardite *et al.*, 1998; Rogler *et al.*, 1998; Schreiber *et al.*, 1998). Activation is evident in the inflamed mucosa but not in uninflamed mucosa (Ardite *et al.*, 1998; Rogler *et al.*, 1998) and is associated with increased IL-8 mRNA expression in the same sites (Ardite *et al.*, 1998). Furthermore, corticosteroid treatment strongly inhibits intestinal NF-κB activation and reduces colonic inflammation (Ardite *et al.*, 1998; Schreiber *et al.*, 1998). Again, inhibition of IL-8 production through the inhibition of NF-κB, as has been demonstrated by these compounds would be predicted be beneficial in inflammatory bowel disease.

Animal models of gastrointestinal inflammation provide further support for NF-κB as a key regulator of colonic inflammation. Increased NF-κB activity is observed in the lamina propria macrophages in 2,4,6-trinitrobenzene sulfonic acid (TNBS)-induced colitis in mice with p65 being a major component of the activated complexes (Neurath *et al.*, 1996; Neurath and Petersson, 1997). Local administration of p65 antisense abrogates the signs of established colitis in the treated animals with no signs of toxicity (Neurath *et al.*, 1996; Neurath and Petersson, 1997). As such, one would predict that small molecule inhibitors of NF-κB would be useful in the treatment of IBD.

Further evidence for a role of NF-κB in inflammatory disorders comes from studies of rheumatoid synovium. Although NF-κB is normally present as an inactive cytoplasmic complex, recent immunohistochemical studies have indicated that NF-κB is present in the nuclei, and hence active, in the cells comprising human rheumatoid synovium (Handel *et al.*, 1995; Marok *et al.*, 1996; Sioud *et al.*, 1998) and in animal models of the disease (Tsao *et al.*, 1997). The staining is associated with type A synoviocytes and vascular endothelium (Marok *et al.*, 1996). Furthermore, constitutive activation of NF-κB is seen in cultured synoviocytes (Roshak *et al.*, 1996; Miyazawa *et al.*, 1998) and in synovial cell cultures stimulated with IL-1β or TNFα (Roshak *et al.*, 1996; Fujisawa *et al.*, 1996; Roshak *et al.*, 1997). Thus, the activation of NF-κB may underlie the increased cytokine production and leukocyte infiltration characteristic of inflamed synovium. The ability of these compounds to inhibit NF-κB and thereby inhibit the production of eicosanoids by these cells would be predicted to yield benefit in rheumatoid arthritis (*see* Table 2).



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Corporation, Madison, WI) into each well and the integrated light output is recorded for 20 sec. Light output is measured in relative light units (RLUs).

NF- $\kappa$ B activity may also be measured in an electrophoretic mobility shift assay

(EMSA) to assess the presence of NF- $\kappa$ B protein in the nucleus. The cells of interest are cultured to a density of  $1 \times 10^6$ /ml. The cells are harvested by centrifugation, washed in PBS without  $\text{Ca}^{2+}$  and  $\text{Mg}^{2+}$ , and resuspended in PBS with  $\text{Ca}^{2+}$  and  $\text{Mg}^{2+}$  at  $1 \times 10^6$  cells/ml.

To examine the effect of compound on the activation of NF- $\kappa$ B, the cell suspensions are treated with various concentrations of drug or vehicle (DMSO, 0.1%) for 30 min at  $37^\circ\text{C}$  prior to stimulation with TNF $\alpha$  (5.0ng/ml) for an additional 15 min. Cellular and nuclear

extracts are prepared follows. Briefly, at the end of the incubation period the cells ( $1 \times 10^6$  cells) are washed 2x in PBS without  $\text{Ca}^{2+}$  and  $\text{Mg}^{2+}$ . The resulting cell pellets are

resuspended in 20  $\mu$ l of Buffer A (10mM Hepes (pH 7.9), 10mM KCl, 1.5mM  $\text{MgCl}_2$ , 0.5mM dithiothreitol (DTT) and 0.1% NP-40) and incubated on ice for 10 min. The nuclei are pelleted by microcentrifugation at 3500 rpm for 10 min at  $4^\circ\text{C}$ . The resulting

supernatant was collected as the cellular extract and the nuclear pellet was resuspended in 15  $\mu$ l Buffer C (20mM Hepes (pH 7.9), 0.42M NaCl, 1.5mM  $\text{MgCl}_2$ , 25% glycerol, 0.2mM EDTA, 0.5mM DTT, and 0.5mM phenylmethylsulphonyl fluoride (PMSF)). The

suspensions are mixed gently for 20 min at  $4^\circ\text{C}$  then microcentrifuged at 14,000 rpm for 10 min at  $4^\circ\text{C}$ . The supernatant is collected and diluted to 60  $\mu$ l with Buffer D (20mM Hepes (pH 7.9), 50mM KCl, 20% glycerol, 0.2mM EDTA, 0.5mM DTT, and 0.5mM PMSF). All samples are stored at  $-80^\circ\text{C}$  until analyzed. The protein concentration of the extracts is determined according to the method of Bradford (Bradford, 1976) with BioRad reagents.

The effect of compounds on transcription factor activation is assessed in an electrophoretic mobility shift assay (EMSA) using nuclear extracts from treated cells as described above. The double stranded NF- $\kappa$ B consensus oligonucleotides ( $5'$ -

AGTTGAGGGGAGCTTCCAGGC- $3'$ ) are labelled with T $_3$  polynucleotide kinase and ( $\gamma$ - $^{32}\text{P}$ )ATP. The binding mixture (25  $\mu$ l) contains 10mM Hepes-NaOH (pH 7.9), 4mM Tris-HCl (pH 7.9), 60mM KCl, 1mM EDTA, 1mM dithiothreitol, 10% glycerol, 0.3 mg/ml bovine serum albumin, and 1  $\mu$ g poly(dI-dC) $\cdot$ poly(dI-dC). The binding mixtures (10  $\mu$ g nuclear extract protein) are incubated for 20 min at room temperature with 0.5 ng of  $^{32}\text{P}$ -labelled oligonucleotide (50,000-100,000 cpm) in the presence or absence of unlabeled competitor after which the mixture is loaded on a 4% polyacrylamide gel prepared in 1X

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#### Biological Assays

The compounds of this invention may be tested in one of several biological assays to determine the concentration of compound which is required to have a given pharmacological effect.

Assays of NF- $\kappa$ B activity are conducted using a cell based luciferase reporter assay as described in Breton, J. J. and Chabot-Fletcher, M. C., *JPET*, 282, 459-466 (1997).

Briefly, U937 human histiocyte lymphoma cell line permanently transfected with the NF- $\kappa$ B reporter plasmids (see below) are cultured in the above medium with the addition of 250  $\mu$ g/ml Geneticin (G418 sulfate, Life Technologies, Grand Island, NY). The luciferase reporter assay is conducted in the transfected U937 clones. These are twice centrifuged at 300 xg for 5 min and resuspended in RPMI 1640 with 10% FBS to a density of  $1 \times 10^6$  cells/ml. One ml aliquots are added to the wells of 24-well plates. Compound or dimethyl

sulfoxide (DMSO) carrier (1  $\mu$ l) is added to the appropriate wells and the plates are incubated at  $37^\circ\text{C}$ , 5%  $\text{CO}_2$  for 30 min. The stimulus is added (5 ng/ml TNF $\alpha$ , 100 ng/ml

LPS, or 0.1  $\mu$ M PMA) and the samples incubated for 5 hours at  $37^\circ\text{C}$ , 5%  $\text{CO}_2$ , transferred to 1.9 ml polypropylene tubes, and centrifuged at 200 xg for 5 min. The cell pellets are washed twice in 1 ml PBS without  $\text{Ca}^{2+}$  and  $\text{Mg}^{2+}$ , and centrifuged as indicated above. The resulting cell pellets are lysed in 50  $\mu$ l 1x lysis buffer (Promega Corporation, Madison, WI), vortexed and incubated for 15 min at room temperature. A 20  $\mu$ l aliquot of each lysate is transferred to an opaque white 96-well plate (Wallac Inc., Gaithersburg, MD) and assayed

for luciferase production in a MicroLumat LB 96 P luminometer (EG&G Berthold, Bad Wilbad, Germany). The luminometer dispenses 100  $\mu$ l luciferase assay reagent (Promega

untreated ears. The inflamed left ears are then removed and stored at -70° until assayed for myeloperoxidase (MPO) activity, a measure of inflammatory cell infiltration.

Inflammatory cell infiltration is assessed through the measurement of

myeloperoxidase activity present in the inflamed ear tissue. Partially thawed ear tissues are minced and then homogenized (10% w/v) with a Tissumizer homogenizer (Tekmar Co., Cincinnati, OH) in 50 mM phosphate buffer (pH 6) containing 0.5% HTAB. The tissue homogenates are taken through three cycles of freeze-thaw, followed by brief sonication (10s). MPO activity in the homogenates is determined as follows. The appearance of colored product from the MPO-dependent reaction of o-dianisidine (0.167 mg/ml, Sigma Chemical, St. Louis, MO) and hydrogen peroxide (0.0005%) is measured spectrophotometrically at 460 nm. Supernatant MPO activity is quantified kinetically (change in absorbance measured over 3 min, sampled at 15 s intervals) using a Beckman DU-7 spectrophotometer and a kinetics analysis package (Beckman Instruments, Inc., Somers, NJ). One unit of MPO activity is defined as that degrading one micromole of

peroxide per minute at 25°C.

Effects on inflammation-mediated cartilage breakdown is measured in an *in vitro* cartilage explant system. In this model bovine articular cartilage explants are incubated for 4 days/96 hours with or without rHuIL-1 alpha to stimulate cartilage breakdown in the presence or absence of test compound. The supernatants are removed for the nitric oxide assays. Nitric oxide was measured using the Greiss reaction and read spectrophotometrically at 530nm. This reaction measures nitrite (NO<sub>2</sub>) which is the stable end product of nitric oxide.

#### General

Nuclear magnetic resonance spectra were recorded at either 250, 300 or 400 MHz using, respectively, a Bruker AM 250, Bruker ARX 300 or Bruker AC 400 spectrometer.

CDCl<sub>3</sub> is deuteriochloroform, DMSO-d<sub>6</sub> is hexadeuteriodimethylsulfoxide, and CD<sub>3</sub>OD is tetra-deuterioethanol. Chemical shifts are reported in parts per million (δ) downfield from the internal standard tetramethylsilane. Abbreviations for NMR data are as follows: s = singlet, d = doublet, t = triplet, q = quartet, m = multiplet, dd = doublet of doublets, dt = doublet of triplets, app = apparent, br = broad. J indicates the NMR coupling constant measured in Hertz. Continuous wave infrared (IR) spectra were recorded on a Perkin-

Elmer 683 infrared spectrometer, and Fourier transform infrared (FTIR) spectra were

Tris borate/EDTA and electrophoresed at 200 V for 2 h. Following electrophoresis the gels are dried and exposed to film for detection of the binding reaction.

The effect of compounds on the phosphorylation of IκB may be monitored in a

Western blot. Cellular extracts are subjected to sodium dodecyl sulfate-polyacrylamide gel electrophoresis (SDS-PAGE) on 10% gels (BioRad, Hercules, CA) and the proteins transferred to nitrocellulose sheets (Hybond™-ECL, Amersham Corp., Arlington Heights, IL). Immunoblot assays are performed using a polyclonal rabbit antibody directed against IκBα or IκBβ followed with a peroxidase-conjugated donkey anti-rabbit secondary antibody (Amersham Corp., Arlington Heights, IL). Immunoreactive bands are detected using the Enhanced Chemiluminescence (ECL) assay system (Amersham Corp., Arlington Heights, IL).

Effects on eicosanoid production by human synovial fibroblasts (RSF) are assessed

using primary cultures of human RSF. These are obtained by enzymatic digestion of

synovium obtained from adult patients with rheumatoid. Cells are cultured in Earl's

Minimal Essential Medium (EMEM) containing 10% fetal bovine serum (FBS), 100

units/ml penicillin and 100 µg/ml streptomycin (GIBCO, Grand Island, NY), at 37°C and

5% CO<sub>2</sub>. Cultures are used at passages 4 through 9 in order to obtain a more uniform type

I fibroblast population. For some studies, fibroblasts are plated at 5 x 10<sup>4</sup> cells/ml in

16mm (diameter) 24 well plates (Costar, Cambridge, MA). Cells are exposed to an optimal

dose of IL-1β (1ng/ml; Roshak *et al.* 1996a)(Genzyme, Cambridge, MA) for the designated

time. Drugs in DMSO vehicle (1%) are added to the cell cultures 15 minutes prior to the

addition of IL-1. Prostaglandin E<sub>2</sub> levels in cell-free medium collected at the termination of

the culture period are directly measured using enzyme immunoassay (EIA) kits purchased

from Cayman Chemical Co. (Ann Arbor, MI). Sample or standard dilutions are made with

experimental medium.

Anti-inflammatory activity *in vivo* is assessed using the phorbol ester-induced ear

inflammation model in mice. Phorbol myristate acetate (PMA) (4µg/20µl acetone) is

applied to the inner and outer surfaces of the left ear of Male Balb/c mice (6/group)

(Charles River Breeding Laboratories, Wilmington, MA). Four hours later, compound

dissolved in 25µl acetone is applied to the same ear. The thickness of both ears is

measured with a dial micrometer (Mitutoyo, Japan) after 20 hours and a second topical

dose of compound is applied. Twenty-four hours later, ear thickness measurements are

taken and the data expressed as the change in thickness (x 10<sup>-3</sup>cm) between treated and

recorded on a Nicolet Impact 400 D infrared spectrometer. IR and FTIR spectra were recorded in transmission mode, and band positions are reported in inverse wavenumbers ( $\text{cm}^{-1}$ ). Mass spectra were taken on either VG 70 FE, PE Syx API III, or VG ZAB HF instruments, using fast atom bombardment (FAB) or electrospray (ES) ionization techniques. Elemental analyses were obtained using a Perkin-Elmer 240C elemental analyzer. Melting points were taken on a Thomas-Hoover melting point apparatus and are uncorrected. All temperatures are reported in degrees Celsius.

Anattech Silica Gel GF and E. Merck Silica Gel 60 F-254 thin layer plates were used for thin layer chromatography. Both flash and gravity chromatography were carried out on E. Merck Kieselgel 60 (230-400 mesh) silica gel.

Where indicated, certain of the materials were purchased from the Aldrich Chemical Co., Milwaukee, Wisconsin, TCI America, Portland, OR.

### Examples

In the following synthetic examples, temperature is in degrees Centigrade ( $^{\circ}\text{C}$ ). Unless otherwise indicated, all of the starting materials were obtained from commercial sources. Without further elaboration, it is believed that one skilled in the art can, using the preceding description, utilize the present invention to its fullest extent. These Examples are given to illustrate the invention, not to limit its scope. Reference is made to the claims for what is reserved to the inventors hereunder.

### Example 1

#### Preparation of N-(4-Acetylphenyl)-2-hydroxy-5-iodobenzoamide.

A solution of iodosalicylic acid (1.9g, 7.4 mmol) and 4-aminoacetophenone (0.97g, 7.4 mmol) in chlorobenzene (40 mL) was treated with  $\text{PCl}_3$  (0.323 mL, 3.7 mmol). The solution was heated at reflux under an argon atmosphere. After 2 h, the solution was filtered hot and the filtrate was left standing at RT. After 18 h solution was filtered and the solid was recrystallized from MeOH to give the title compound (0.095 g, 5% yield):  $^1\text{H}$  NMR (400 MHz,  $\text{DMSO}-d_6$ )  $\delta$  2.5-2.6 (s, 3H), 6.8-8.2 (m, 7H), 10.1-10.2 (s, 1H).

### Example 2

#### Preparation of N-(2,4-Difluorophenyl)-2-hydroxy-5-nitrophenylcarboxamide

A solution of 5-nitrosalicylic acid (1.4 g, 7.7 mmol) and 2,4-difluoroaniline (0.8 mL, 7.7 mmol) in chlorobenzene (40 mL) was treated with  $\text{PCl}_3$  (0.338 mL, 3.8 mmol). The solution was heated at reflux under an argon atmosphere. After 2 h, the solution was filtered hot and the filtrate was left standing at RT. After 18 h the solution was filtered and the solid was recrystallized from MeOH to give the title compound (0.733g, 35% yield):  $^1\text{H}$  NMR (400 MHz,  $\text{DMSO}-d_6$ )  $\delta$  2.5-2.6 (s, 3H), 7.1-8.9 (m, 6H), 10.6-10.7 (s, 1H)

### Example 3

#### Preparation of N-(2,4-Difluorophenyl)-2-hydroxy-5-iodophenylcarboxamide

a) 5-Iodosalicylic acidchloride

5-Iodosalicylic acid (2.0 g, 7.38 mmol) in toluene was treated with  $\text{SOCl}_2$  (1.66 mL, 22.7 mmol) and catalytic DMF at reflux for 1 h. The reaction mixture was evaporated to dryness and the acid chloride used in the next step without purification.

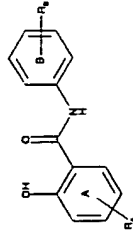
b) 2,4-Difluorophenyl)-2-hydroxy-5-iodophenylcarboxamide

The compound of Example 3(a) (3.79 mmol) and 2,4-difluoroaniline (380  $\mu\text{L}$ , 3.79 mmol) in toluene was heated at reflux for 24 h. The reaction mixture was evaporated, the residue washed with ether and the solid residue recrystallized from MeOH to give 159 mg of N-(2,4-difluorophenyl)-2-hydroxy-5-iodophenylcarboxamide. ES MS ( $\text{M}+\text{H}$ ) $^{+}$  m/e 373.7.

The above specification and Examples fully disclose how to make and use the compounds of the present invention. However, the present invention is not limited to the particular embodiments described hereinabove, but includes all modifications thereof within the scope of the following claims. The various references to journals, patents and other publications which are cited herein comprise the state of the art and are incorporated herein by reference as though fully set forth.

We claim:

1. A pharmaceutical composition comprising a compound of Formula I:



I

wherein:

- $R_A$  substitutes ring A 0-3 times and is independently selected from the group consisting of:  $\text{NO}_2$ , halogen,  $\text{C}_1$ -alkyl, trifluoromethyl, O- $\text{C}_1$ -alkyl and S- $\text{C}_1$ -alkyl; and
- $R_B$  substitutes ring B 0-3 times and is independently selected from the group consisting of: halogen,  $\text{C}(\text{O})\text{C}_1$ -alkyl,  $\text{C}_1$ -alkyl, O- $\text{C}_1$ -alkyl, S- $\text{C}_1$ -alkyl,  $\text{CH}_2$ -aryl, and aryl; and
- and pharmaceutically acceptable salts, hydrates and solvates thereof.

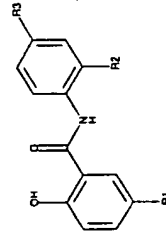
2. A pharmaceutical composition according to Claim 1 wherein:

$R_A$  occurs once and is  $\text{R}_1$ ; and

$R_B$  occurs twice and is independently  $\text{R}_2$  and  $\text{R}_3$ .

3. A pharmaceutical composition according to Claim 1 comprising a compound of

Formula II:



II

wherein:

- $\text{R}_1$  is selected from the group consisting of: H,  $\text{NO}_2$ ,  $\text{CF}_3$ , F, Cl, Br, and I;
- $\text{R}_2$  is selected from the group consisting of: H and F; and

$\text{R}_3$  is selected from the group consisting of: F, Cl, Br, I, phenyl and  $\text{C}(\text{O})\text{C}_1$ -alkyl;

and a pharmaceutically acceptable carrier, diluent or excipient.

4. A pharmaceutical composition according to Claim 3 wherein  $\text{C}(\text{O})\text{C}_1$ -alkyl is  $\text{C}(\text{O})\text{CH}_3$ .

5. A pharmaceutical composition according to Claim 3 wherein said compound is selected from the group consisting of:

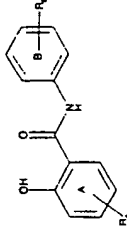
N-(4-phenyl-phenyl)-2-hydroxy-5-trifluoromethylcarboxamide;

N-(2,4-Difluorophenyl)-2-hydroxy-5-nitrocarboxamide;

N-(2,4-Difluorophenyl)-2-hydroxy-5-iodocarboxamide; and

N-(4-acetylphenyl)-2-hydroxy-5-iodocarboxamide.

6. A method of inhibiting NF- $\kappa\text{B}$  comprising administering to a patient in need thereof an effective amount of a compound of Formula I:



I

wherein:

- $R_A$  substitutes ring A 0-3 times and is independently selected from the group consisting of:  $\text{NO}_2$ , halogen,  $\text{C}_1$ -alkyl, trifluoromethyl, O- $\text{C}_1$ -alkyl and S- $\text{C}_1$ -alkyl; and

$R_B$  substitutes ring B 0-3 times and is independently selected from the group consisting of: halogen,  $\text{C}(\text{O})\text{C}_1$ -alkyl,  $\text{C}_1$ -alkyl, O- $\text{C}_1$ -alkyl, S- $\text{C}_1$ -alkyl,  $\text{CH}_2$ -aryl, and aryl;

- and pharmaceutically acceptable salts, hydrates and solvates thereof.

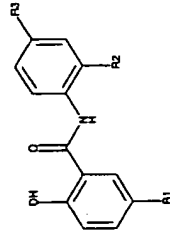
7. A method according to Claim 6 wherein:

$R_A$  occurs once and is  $R_1$ ; and

$R_B$  occurs twice and is independently  $R_2$  and  $R_3$ .

5

8. A method according to Claim 7 comprising administering to a patient in need thereof an effective amount of a compound of Formula II:



10

wherein:

$R_1$  is selected from the group consisting of: H,  $NO_2$ ,  $CF_3$ , F, Cl, Br, and I;

$R_2$  is selected from the group consisting of: H and F; and

$R_3$  is selected from the group consisting of: F, Cl, Br, I, phenyl and

$C(O)C_1-6alkyl$ .

15

9. A method according to Claim 8 wherein said compound is selected from the group consisting of:

N-(4-phenyl-phenyl)-2-hydroxy-5-trifluoromethylcarboxamide;

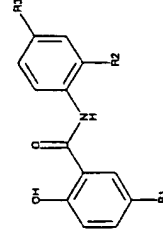
N-(2,4-Difluorophenyl)-2-hydroxy-5-nitrocarboxamide;

N-(2,4-Difluorophenyl)-2-hydroxy-5-iodocarboxamide; and

N-(4-acetylphenyl)-2-hydroxy-5-iodocarboxamide.

20

12. A method according to Claim 11 comprising administering to a patient in need thereof an effective amount of a compound of Formula I:



11

wherein:

$R_1$  is selected from the group consisting of: H,  $NO_2$ ,  $CF_3$ , F, Cl, Br, and I;

$R_2$  is selected from the group consisting of: H and F; and

25

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R<sub>3</sub> is selected from the group consisting of: F, Cl, Br, I, phenyl and C(O)C1-6alkyl.

13. A method according to Claim 12 wherein said compound is selected from the group  
5 consisting of:

N-(4-phenyl-phenyl)-2-hydroxy-5-trifluoromethylcarboxamide;

N-(2,4-Difluorophenyl)-2-hydroxy-5-nitrocarboxamide;

N-(2,4-Difluorophenyl)-2-hydroxy-5-iodocarboxamide; and

N-(4-acetylphenyl)-2-hydroxy-5-iodocarboxamide.

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14. A method according to Claims 10 to 13 wherein said disease is an inflammatory  
disorder.

15. A method according to Claim 14 wherein said disease is selected from the group  
15 consisting of: rheumatoid arthritis, inflammatory bowel disease, and asthma.

16. A method according to Claim 10 to 13 wherein said disease is dermatosis.

17. A method according to Claim 16 wherein said disease is selected from the group  
20 consisting of: psoriasis and atopic dermatitis.

18. A method according to Claim 10 to 13 wherein said disease is selected from the  
group consisting of: autoimmune diseases; tissue and organ rejection; Alzheimer's disease;  
stroke; atherosclerosis; stenosis; osteoarthritis; osteoporosis; and Ataxia Telangiectasia.

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19. A method according to Claim 10 to 13 wherein said disease is cancer.

20. A method according to Claim 19 wherein said cancer is Hodgkins disease.

21. A method according to Claim 10 to 13 wherein said disease is AIDS.

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